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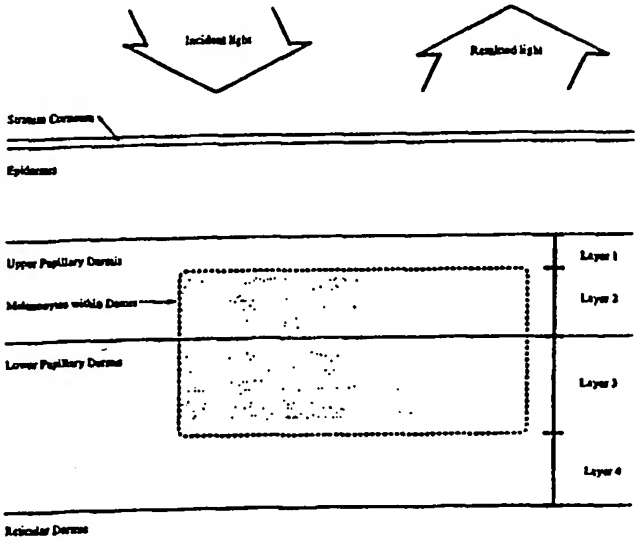

(57) Abstract

The invention relates to a method for non-invasively measuring skin structure. Infrared radiation from a plurality of locations over an area of skin under investigation is measured so as to give an indication of the variation in papillary dermis thickness over said area, and the skin colour coordinates at a plurality of locations over the same area of skin is also measured. The data obtained is used to calculate corrected skin colour coordinates over the area corresponding to a predetermined papillary dermis thickness. The corrected skin colour coordinates so obtained are compared with a reference colour coordinate range for healthy skin of the same predetermined papillary dermis thickness. At an abnormal region, where the corrected skin colour coordinates lie outside the reference colour coordinate range, the depth of penetration of dermal melanin can be measured.

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METHOD FOR MEASUREMENTOF SKIN HISTOLOGY

This invention relates to a method for the non-invasive measurement of skin histology and is particularly, but not exclusively, concerned with a method for identifying and measuring the presence and depth of dermal invasion of melanin. The presence and extent of dermal invasion within a skin cancer is considered to be the most important factor governing a patient's prognosis. The present invention is considered to be potentially useful for the preliminary screening of patients to identify those who should be referred to an appropriate clinician for diagnosis and further to assist the clinician in diagnosis.

The present invention is based on the findings reported by Symon D'O Cotton in "Do all human skin colours lie on a defined surface within LMS space?", University of Birmingham Technical Report, 30 December 1995. The disclosure of such Technical Report is included herein by reference. In this Technical Report, the relation between healthy skin and the colour of the skin represented in LMS, a particular colour space, is reported, and it discloses that, for healthy skin, the coloration, regardless of race or amount of tanning, lies on a defined curved surface within a three-dimensional colour space. This, if used with a correct colour measurement system, can measure and quantify the amount of melanin and blood at any particular point at which this measurement is made. If the skin is sampled as an image, then corresponding images showing the variation of blood and melanin across the skin can be obtained. In the above Technical Report, it is disclosed that melanin can sometimes penetrate into the dermis producing the characteristic hues of melanoma and that this melanocytic descent has been quantified by Clark et al ("The

Histogenesis and Biological Behaviour of Primary Human Malignant Melanomas of the Skin", Cancer Research, 29, 1989) into five levels of tumour invasion, in which level 1 corresponds to confinement within the epidermis, level 2 corresponds to invasion into the papillary dermis, etc. In an alternative system, the extent of tumour invasion in mm from the cornified layer is expressed as the Breslow thickness. The above Technical Report also acknowledges that, in the case of melanoma, CD Neville ("Melanoma: Issues of Importance to the Clinician", British Journal of Hospital Medicine, March 1985) discloses the existence of a strong relationship between this level of invasion and prognosis. However, the above Technical Report does not disclose in detail any method suitable for taking the necessary measurements.

According to the present invention, there is provided a method of non-invasively analysing skin structure, comprising the steps of:
(i) measuring infrared radiation from a plurality of locations over an area of skin under investigation so as to give an indication of the variation in papillary dermis thickness over said area;
(ii) measuring the skin colour coordinates at a plurality of locations over said area of skin;
(iii) using data obtained in measuring steps (i) and (ii) to calculate corrected skin colour coordinates over said area which corresponds to a predetermined papillary dermis thickness, and;
(iv) comparing the corrected skin colour coordinates obtained in step (iii) with a reference colour coordinate range for healthy skin of the same predetermined papillary dermis thickness.
The method can be used for locating and measuring the properties of a skin abnormality, in which case the method further comprises the steps of;

(v) identifying an abnormal location (i.e. a region where melanin exists within the dermis) within said area of skin where the corrected skin colour coordinates lie outside the reference colour coordinate range;
(vi) calibrating the corrected skin colour coordinates of said abnormal location with the corrected skin colour coordinates of at least one skin location having colour coordinates lying within said reference colour coordinate range for normal skin, and;
(vii) using the skin colour coordinates to assess the degree of abnormality of said abnormal skin location.

It is to be understood that using this method, it is possible to reconstruct a full 3D model of the skin architecture which conveys information grossly comparable to that available through microscopical examination of biopsied skin tissue.

It has been found that the papillary dermal skin thickness can change markedly with some skin lesions which are not otherwise of concern. This throws the coloration of the skin off the surface of predicted coloration and so can give rise to false measurements of the histology of such skin lesions. It is for this reason that papillary dermis thickness is measured first, and subsequent calculations are based on the skin colour coordinates corrected to a predetermined papillary dermis thickness. Any arbitrary value for this thickness may be chosen, such as 2.0×10^{-4} m which is the average value for healthy human skin.

The thickness of the papillary dermis may be obtained by utilising the property of human skin to vary its absorption of infrared radiation with varying papillary dermis thickness. In general, there is an inverse

relationship between absorption and thickness. The fact that infrared radiation is also absorbed by other materials within the skin, particularly melanin and blood, is a complicating factor. However the effect on absorption of varying blood and melanin content is far smaller than the effect of papillary dermis thickness, and so the latter may still be measured. This can be done by obtaining two infrared images, each at a different wavelength. The chosen wavelengths are not important, but one should be further into the infrared (ie at longer wavelength) than the other. Suitable wavelength bands are 800-1000nm and 600-800nm, in that readily available infrared films and filters may be used. The brightness of points within the image obtained at the longer wavelength is affected to a greater extent by variations in the papillary dermis thickness. Conversely, the image obtained at shorter wavelength will be affected to a greater extent by other materials such as melanin and blood. By predicting the brightnesses of points of differing papillary dermis thickness and amounts of epidermal melanin which absorb near-infrared radiation at the two different infrared wavelengths, a reference graph (Fig 1) can be obtained which consists of lines of constant papillary dermis thickness, wherein Primary 1 is the measurement made at the longer (800-1000nm) wavelength and Primary 2 is the measurement made at the shorter (600-800nm) wavelength. The absorption of blood within these wavelengths is very small (a hundredth of its peak value for visible wavelengths at 600-800nm and even less for 800-1000nm) and to a first approximation may be ignored. The presence of dermal melanin does introduce a small error in the range of low values for both primaries, but this is insignificant in practice. Thus, by comparing values obtained at these wavelengths with this graph, it is possible to ascertain the papillary dermis thickness. However it is within the scope of the present invention to measure

brightness at such a long infra-red wavelength eg. 1100nm that the brightness would vary to such a negligible extent with melanin and blood content that it would effectively depend solely on the papillary dermis thickness. This would also reduce the error introduced by the presence of dermal melanin. In such a case only one set of brightness measurements would be required. Furthermore, a transformation can be calculated which allows an image of the skin to be created which represents how the skin would appear if it had a papillary dermis thickness of any predetermined value.

In a preferred embodiment, the reference colour coordinate range for normal skin at the predetermined papillary dermis thickness is obtained as disclosed in the above-mentioned Technical Report as a curved surface lying within a three-dimensional colour space, with one of the bounding axes relating to the amount of melanin within the epidermis and the other relating to the amount of blood within the dermis. When an area containing dermal melanin is located, i.e. points do not lie on the normal colour surface, the epidermal melanin value within this area is estimated by either reference to the reference colour coordinate range for normal skin within regions identified as normal, or by reference to the epidermal melanin levels calculated within normal regions adjacent to said area containing dermal melanin. This value is then used with the corrected colour coordinates of the abnormal region at the same predetermined papillary dermis thickness to compute invasion depth and concentration of dermal melanin. The corrected skin colour coordinates for the area of skin under investigation may be calibrated to values equivalent to zero epidermal melanin. Instead of using LMS colour space, it is possible to

use any other colour space, for example, the RGB colour space or a UV G IR colour space.

The dermis contrasts strongly in structure to that of the epidermis, being highly vascular, containing many sensory receptors and being made largely from collagen fibres to provide the essential structure of the skin. Between the epidermis and the dermis, the junction presents an extremely uneven boundary with finger-like dermal protrusions called dermal papillae projecting towards the skin surface. The dermis can be split into two further histologically distinct layers, the *papillary dermis* and the reticular dermis within which the structure of the collagen fibres differs significantly. The papillary dermis is situated directly below the epidermis and within which the collagen exists as a fine network of fibres. This is in contrast with the reticular dermis where the collagen fibres are aggregated into thick bundles which are arranged nearly parallel to the skin surface. In the case of melanin invasion of the papillary dermis, there is a layer containing blood, melanin and collagen, a layer containing either blood and collagen or melanin and collagen, depending upon whether melanin has passed the blood layer; and a layer containing just collagen. The different thicknesses of these layers, the amount of blood and the concentration of dermal melanin along with the amount of melanin in the overlying epidermis affect the remitted light. This can be modelled by calculating the net effect of these three layers for the differing parameters outlined.

A mathematical model describing the optics of the skin has been described in the above mentioned Symon D'O Cotton's Technical Report, whose disclosure has been included herein by reference, and this model

can be extended to predict coloration of skin containing dermal descent of melanin.

As can be seen from Fig 2, there are now four distinct layers within the dermis which can combine to construct a simple model, 1) a layer within the upper papillary dermis containing no melanin, 2) a layer within the upper papillary dermis containing melanin, 3) a layer within the lower papillary dermis containing melanin, 4) a layer within the lower papillary dermis containing no melanin.

It should also be noted that the condition of melanin existing up to the dermo-epidermal junction is facilitated by allowing the thickness of layer 1 to be zero and likewise melanin can exist up to the papillary-reticular dermis boundary by setting the thickness of layer 4 to be zero.

In computing a model to predict this coloration it is useful to make note of the fact that, as discussed in section 2.1 of the Technical Report, the amount of back scatter due to melanin can be considered negligible. Therefore, in the same manner that it was possible to apply the Kubelka-Munk theory to the papillary dermis (section 3.2.2 of the Technical Report), to compute the coloration of sections of papillary dermis containing blood, where the back scattering component of blood was considered negligible, it is possible to compute the coloration of sections containing melanin. In this situation $\varsigma(\lambda)$ (scattering coefficient) remains dependent only on wavelength whilst α (fraction of radiation absorbed per unit path length) becomes $\alpha(\lambda,\rho,\Phi)$ where Φ represents the density of dermal melanin within that layer. Further, following the proof given in equation (17) of the Technical Report, $\alpha(\lambda,\rho,\Phi)$ can be shown to be the

sum of $\alpha_{iv}(\lambda)$, $\alpha_b(\lambda)$ and $\alpha_m(\lambda)$, where $\alpha_m(\lambda)$ is the absorption coefficient of melanin. From the above it is possible to calculate R and T (diffuse radiation and transmission respectively). For simplicity of notation it is helpful to consider R_1 & T_1 where,

$$R_1(\lambda,\rho,\Phi,d_n) = R(\beta(k(\alpha(\lambda,\rho,\Phi)),s(\varsigma(\lambda))),K(k(\alpha(\lambda,\rho,\Phi)),s(\varsigma(\lambda))),d_n)$$

and

$$T_1(\lambda,\rho,\Phi,d_n) = T(\beta(k(\alpha(\lambda,\rho,\Phi)),s(\varsigma(\lambda))),K(k(\alpha(\lambda,\rho,\Phi)),s(\varsigma(\lambda))),d_n)$$

where d_n is the layer thickness.

As was shown in section 3.2.3 of the Technical Report, two-layer systems can be combined to produce the total remitted and transmitted light for the dermis resulting in equation (20) of the Technical Report.
This can be simplified using the geometric series

$$a + ar + ar^2 + ar^3 + \cdots = \frac{a}{1-r} \text{ if } -1 < r < 1$$

to

$$R_{1total}(\lambda,\rho_{ud},\rho_{ld},d_{ud},d_{ld}) = R_{1ud}(\lambda,\rho_{ud},d_{ud}) + \frac{T_{1ud}(\lambda,\rho_{ud},d_{ud})^2 R_{1ld}(\lambda,\rho_{ld},d_{ld})}{1 - R_{1ud}(\lambda,\rho_{ud},d_{ud})R_{1ld}(\lambda,\rho_{ld},d_{ld})}$$

Similarly, T_{1total} can be shown to be

$$T_{1total}(\lambda,\rho_{ud},\rho_{ld},d_{ud},d_{ld}) = \frac{T_{1ud}(\lambda,\rho_{ud},d_{ud}) * T_{1ld}(\lambda,\rho_{ld},d_{ld})}{1 - R_{1ud}(\lambda,\rho_{ud},d_{ud})R_{1ld}(\lambda,\rho_{ld},d_{ld})}$$

These equations can be extended, as is shown by Wan et al. [1981], to an n layered system resulting in values for $R_{12...n}$ and $T_{12...n}$ of

$$R_{12...n} = R_{12...n-1} + \frac{T_{12...n-1}^2 R_n}{1 - R_{12...n-1}R_n}$$

$$T_{12...n} = \frac{T_{12...n-1}T_n}{1 - R_{12...n-1}R_n}$$

This system of equations can therefore compute the total remitted and transmitted light from an *n* layered system of arbitrary complexity provided that the thickness and composition of the layers is specified.

For the four-layer system shown in Fig 2, this results in a value for the total light remitted and transmitted from the dermis dependent on λ, P_{ud}, P_{ld}, d_{ud}, d_{ld}, d_{l2}, Φ_{l2}, d_{l3} and Φ_{l3} where d_{l2} and d_{l3} are the thickness of layers 2 and 3 whilst Φ_{l2} and Φ_{l3} are their corresponding melanin densities. The thickness of layer 1 and layer 2 do not need to be explicitly defined as they are simply d_{ud} - d_{l2} and d_{ld} - d_{l3} respectively; similarly Φ_{l1} and Φ_{l4} are zero by definition. A further simplification is possible if it is assumed that $\Phi_{l2} = \Phi_{l3}$ leading to a single value of Φ for the dermis.

The results of these equations can be combined with the predicted light transmitted by the epidermis in the same manner as that discussed in section 3.3 of the Technical Report, thus leading to the following description of total remitted, S_{rd}, and transmitted S_{td}.

$$S_{rd}(\lambda, \rho_{ud}, \rho_{ld}, d_{ud}, d_{ld}, d_{l2}, d_{l3}, \Phi, d_m) = R_{2total}(\lambda, \rho_{ud}, \rho_{ld}, d_{ud}, d_{ld}, d_{l2}, d_{l3}, \Phi)\theta(\lambda, d_m)^2 S(\lambda)$$

$$S_{td}(\lambda, \rho_{ud}, \rho_{ld}, d_{ud}, d_{ld}, d_{l2}, d_{l3}, \Phi, d_m) = T_{2total}(\lambda, \rho_{ud}, \rho_{ld}, d_{ud}, d_{ld}, d_{l2}, d_{l3}, \Phi)\theta(\lambda, d_m)^2 S(\lambda)$$

These can be used to predict the value of the corresponding LMS primaries

$$L(\rho_{ud},\rho_{ld},d_{ud},d_{ld},d_{12},d_{13},\Phi,d_m) = \int_0^\infty R_{2total}(\lambda,\rho_{ud},\rho_{ld},d_{ud},d_{ld},d_{12},d_{13},\Phi)\theta(\lambda,d_m)^2 S(\lambda)S_L(\lambda)d\lambda$$

$$M(\rho_{ud},\rho_{ld},d_{ud},d_{ld},d_{12},d_{13},\Phi,d_m) = \int_0^\infty R_{2total}(\lambda,\rho_{ud},\rho_{ld},d_{ud},d_{ld},d_{12},d_{13},\Phi)\theta(\lambda,d_m)^2 S(\lambda)S_M(\lambda)d\lambda$$

$$S(\rho_{ud},\rho_{ld},d_{ud},d_{ld},d_{12},d_{13},\Phi,d_m) = \int_0^\infty R_{2total}(\lambda,\rho_{ud},\rho_{ld},d_{ud},d_{ld},d_{12},d_{13},\Phi)\theta(\lambda,d_m)^2 S(\lambda)S_S(\lambda)d\lambda$$

A further generalisation can be made to any primary, P_n, leading to the following equation where S_n defines the spectral response of that primary.

$$P_n(\rho_{ud},\rho_{ld},d_{ud},d_{ld},d_{12},d_{13},\Phi,d_m) = \int_0^\infty R_{2total}(\lambda,\rho_{ud},\rho_{ld},d_{ud},d_{ld},d_{12},d_{13},\Phi)\theta(\lambda,d_m)^2 S(\lambda)S_{P_n}(\lambda)d\lambda$$

This equation can then be used to generate the expected coloration of human skin exhibiting dermal descent of melanin.

The result of this analysis is that it is possible for the same coloration to result from different combinations of the above parameters. This complicates the measurement of the dermal invasion of melanin, (but not identifying the presence of any dermal melanin). Indeed, to obtain this measurement, it is necessary to know the amount of melanin in the overlying epidermis. However, at points where dermal invasion has taken place, this parameter is difficult to determine simply by comparing colour coordinates of the abnormal location with colour coordinates for healthy skin. It is for this reason that, in the present invention, regions where dermal melanin exists are identified by reference to a reference colour coordinate range for healthy skin, and then the colour coordinates of these regions are compared with the colour coordinates at one or more normal

skin locations. If said normal skin locations are adjacent to the region where dermal melanin exists, it is sufficient to use the epidermal melanin levels calculated for such normal skin locations to estimate the epidermal melanin levels at the region where dermal melanin exists Alternatively, it is possible to perform a measurement of the epidermal melanin levels within areas of the skin where the presence of dermal melanin has been identified, by assessing the deviation in coloration at the blue end of the spectrum, from the reference colour coordinate range for normal skin due to the presence of such dermal melanin. At the blue end of the spectrum, the increase in such deviation quickly slows with increasing depth of melanin penetration until a "saturation point" is reached. By assuming that the depth of melanin penetration within the dermis is large enough for such saturation to have occurred, an estimate of the deviation from the reference colour coordinate range for normal skin can be made. This estimate allows a calculation to be made of the skin coloration assuming no dermal melanin, and therefore by reference to the colour coordinate range for normal skin, of the level of epidermal melanin. It is within the scope of the present invention to measure the epidermal melanin levels directly, for example using polarised light, and to incorporate such measurements in the measuring step (ii) above.

By any of the above methods, the effect of what would have been the normal epidermal melanin level in the abnormal skin location can be taken into account, thereby enabling a more accurate determination of melanin descent

By comparing the values of the skin image represented in a certain colour space with theoretically calculated values covering all possible amounts of

blood, dermal melanin penetration and melanin concentration within the same colour space, the values of those three parameters can be obtained for every point in the image. Since the papillary dermis thickness and epidermal melanin content are known, it is possible to compute a detailed three-dimensional reconstruction of the top layers of human skin. This is of great potential interest to the medical profession and enables routine examination of the internal structure of living skin just as X-rays, NMR and ultrasound are used for examining other parts of the body. It is also within the scope of the invention to acquire the infra-red and/ or visible images using lasers of different wavelengths or by using spectral analysis.

It is possible to use a computer programmed with the above algorithms to perform the actual calculations. However, before these calculations can be performed, an image of the area of skin under investigation must be represented in the same colour space as for the normal skin reference colour coordinate range. This can be done in a number of ways. In one way, the skin colour coordinates are acquired from an image using the same lighting conditions and a CCD camera calibrated in the same way as that used to produce the healthy skin reference colour coordinate range. Alternatively, if exactly the same lighting conditions are not used, a white standard or other appropriate correction factor can be used to allow calibration of the image within the software. As a further alternative, a colour image can be acquired using a colour photographic film which is then digitised. This can be performed using either exactly the same lighting conditions and a calibrated set-up or again with the inclusion of a white standard or other appropriate correction factor. It is within the scope of this invention to obtain both the infra-red and visible images with

a single digital camera or to calculate the value of the necessary primaries through the use of spectroscopy.

The present invention will now be described in further detail and with reference to the accompanying drawings, in which:-

Fig 1 is a graph showing variation of brightness with papillary dermis thickness for primaries 1 and 2. as described hereinabove;

Fig 2 is a schematic cross-sectional view through a section of skin illustrating melanin descent into the *papillary dermis*;

Fig 3 is a schematic cross-sectional view through a section of skin illustrating normal, healthy regions and an abnormal region where, in this case, melanin descent into the *papillary dermis* and the *reticular dermis* has taken place;

Fig 4 is a block diagram showing the steps involved in one embodiment of the method of the present invention;

Fig 5 is a diagram showing the predicted surface of normal skin coloration within a three-dimensional colour space;

Fig 6 is a diagram showing coloration within the skin cancer that is shown in Fig 7 in the same 3-D colour space as depicted in Fig 5, wherein areas of normal and abnormal coloration are shown; and

Fig 7 is a photographic image of the skin cancer.

Referring now to Fig 3 of the drawings, a schematic skin section is shown wherein melanin (indicated by the black circles in Fig 3) in normal healthy skin are present in the lower part of *epidermis* 10 adjacent but above the dermo-epidermal junction 12 between the *epidermis* and the *papillary dermis* 14. The Breslow thickness referred to above is the depth of melanin invasion in millimetres measured from granular layer 16 which is a layer in the *epidermis* 10 where the skin goes scaly and forms the tough outer cornified layer 18. In the abnormal region of the skin, the melanin is shown as having descended not only into the *papillary dermis* 14, but also into the underlying *reticular dermis* 20 lying above the subcutaneous fat layer 22. It is to be appreciated that, in other cases, melanin decent can be into any layer of the skin and may even be into the subcutaneous fat layer 22.

Referring now to Fig 4, there is shown a block diagram illustrating the steps involved in a typical method of measurement in accordance with the present invention. In Fig 4, block 38 exemplifies method step (i) above- the determination of papillary dermis thickness by shining infrared light at two wavelengths on an area of skin being subjected to measurement and measuring the amount of light reflected from a plurality of points within that area. Block 40 exemplifies method step (ii) above- the acquisition of an image at visible wavelengths of the same skin area. This can be by CCD camera, digitised film or any other convenient means. Block 42 exemplifies method step (iii) above- the transformation of the image into corrected colour space of the skin model at a predetermined papillary dermis thickness. Block 44 exemplifies method steps (iv and v) above- the identification of regions containing dermal melanin, by comparing the

corrected skin colour coordinates with the reference colour coordinate range. Block 46 exemplifies method step (vi) above- use of the corrected colour space to calculate the amounts of *epidermal* melanin within normal regions adjacent to the regions containing dermal melanin and use thereof to give an indication of the amounts thereof which exist in the regions containing dermal melanin. Block 48 exemplifies a first part of method step (vii) above- calculation of *dermal* invasion using the measured coloration of the abnormal regions and the calculated amount of *epidermal* melanin from 46. Block 50 exemplifies a second part of method step (vii) above- transformation of the calculated *dermal* invasion of melanin into either the Breslow thickness or the Clark's level of invasion. This can be reported as either representing the maximum invasion or as an image showing invasion over the skin.

Referring now to Fig 5, the shaded surface indicates the range of colorations which can exist in normal healthy skin corrected to the predetermined papillary dermis thickness. Skin colorations which depart from this surface are indicative of dermal melanin.

Referring now to Figs. 6 and 7, it can be seen that a region of the skin which is shown in Fig 7 and which is indicated by arrow H in Fig 6 lies at a position corresponding to part of the shaded surface illustrated in Fig 5 and is indicative of normal healthy skin, whereas an adjacent region indicated by arrow U in Fig 6 lies outside such surface and is indicative of skin containing dermal melanin. Comparison of the coloration of these two adjacent regions H and U enables the depth of melanin invasion in the abnormal region of the skin in Fig 7 to be computed.

CLAIMS

1. A method of non-invasively analysing skin structure, comprising the steps of:

(i) measuring infrared radiation from a plurality of locations over an area of skin under investigation so as to give an indication of the variation in papillary dermis thickness over said area;

(ii) measuring the skin colour coordinates at a plurality of locations over said area of skin;

(iii) using data obtained in measuring steps (i) and (ii) to calculate corrected skin colour coordinates over said area which corresponds to a predetermined papillary dermis thickness, and;

(iv) comparing the corrected skin colour coordinates obtained in step (iii) with a reference colour coordinate range for healthy skin of the same predetermined papillary dermis thickness.

2. A method according to claim 1, comprising the additional steps of;

(v) identifying an abnormal location within said area of skin where the corrected skin colour coordinates lie outside the reference colour coordinate range;

(vi) calibrating the corrected skin colour coordinates of said abnormal location with the corrected skin colour coordinates of at least one skin location having colour coordinates lying within said reference colour coordinate range for normal skin, and;

(vii) using the skin colour coordinates to assess the degree of abnormality of said abnormal skin location.

3. A method according to claim 1 or 2, wherein an independent measurement of the level of epidermal melanin is made.

4. A method according to claim 3, wherein said independent measurement of the level of epidermal melanin is by polarized light.

5. A method according to claim 2, wherein said calibration in step (vi) includes estimating the level of epidermal melanin in said abnormal location by reference to epidermal melanin levels calculated within at least one normal skin location adjacent said abnormal region.

6. A method according to claim 2, wherein said calibration in step (vi) includes measuring epidermal melanin levels at said abnormal location by assessing the deviation at the blue end of the spectrum at said abnormal location from the reference colour coordinate range for normal skin.

7. A method according to any preceding claim, wherein in step (i), a single infrared image at a wavelength of greater than about 1100nm is obtained for each of said locations.

8. A method according to any one of claims 1 to 6, wherein in step (i) two infrared images, each at a different wavelength, are obtained for each of said locations, whereby to enable the effect of the presence of epidermal melanin and dermal blood to be accounted for in the calculation of step (iii).

9. A method according to claim 8, wherein said wavelengths fall within the bands 800-1000nm and 600-800nm respectively.

10. A method according to any one of claims 7 to 9, wherein said infrared image(s) is/are obtained using infrared photographic film, or laser(s), or by spectral analysis.

11. A method according to any preceding claim, wherein the reference colour coordinate range for normal skin at the predetermined papillary dermis thickness referred to in step (iv), is obtained as a curved surface lying within a three-dimensional colour space, with a first bounding axis relating to the amount of melanin within the epidermis and a second bounding axis relating to the amount of blood within the dermis.

12. A method according to claim 11, wherein said three-dimensional colour space is selected from LMS, RGB and UV G IR colour spaces.

13. A method according to any preceding claim, wherein the skin colour coordinates of step (ii) are acquired from an image using the same lighting conditions and the same calibration set-up as used to produce the healthy skin reference colour coordinate range.

14. A method according to any one of claims 1 to 12, wherein the skin colour coordinates of step (ii) are acquired from an image using different lighting conditions than used to obtain the healthy skin reference colour coordinate range, and a white standard or other correction factor is used to allow calibration of the image with the reference colour coordinate range.

FIG. 1

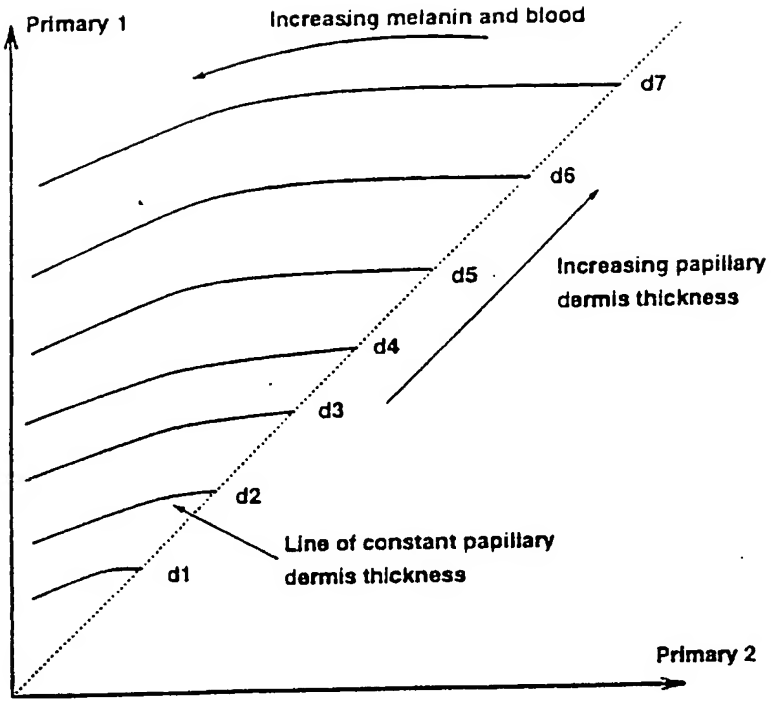
Primary 1
Increasing melanin and blood
d7
d6
d5
d4
d3
d2
d1
Increasing papillary dermis thickness
Line of constant papillary dermis thickness
Primary 2

FIG. 2

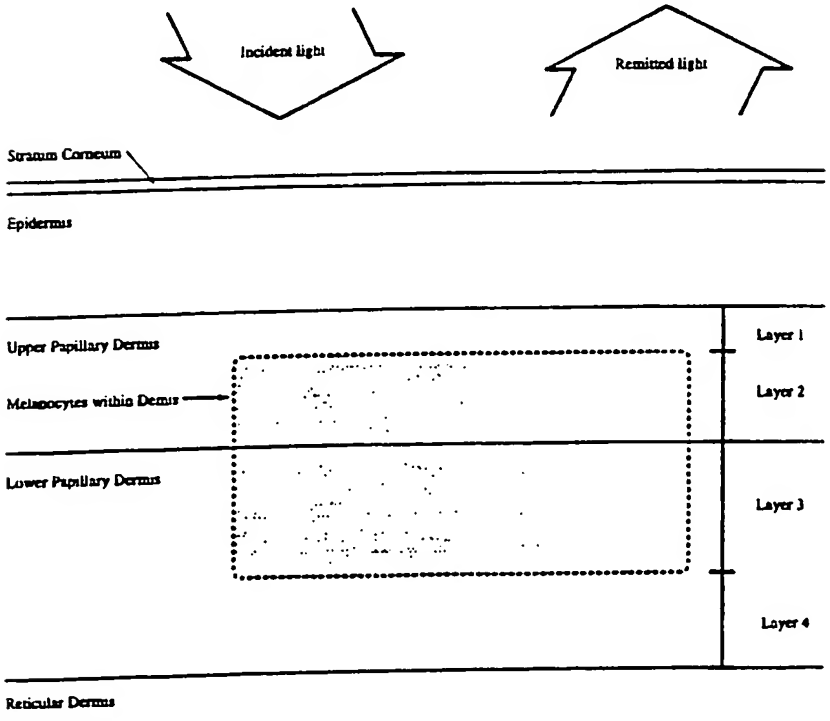
Incident light
Remitted light
Stratum Corneum
Epidermis
Upper Papillary Dermis
Melanocytes within Dermis
Lower Papillary Dermis
Reticular Dermis
Layer 1
Layer 2
Layer 3
Layer 4

FIG. 3

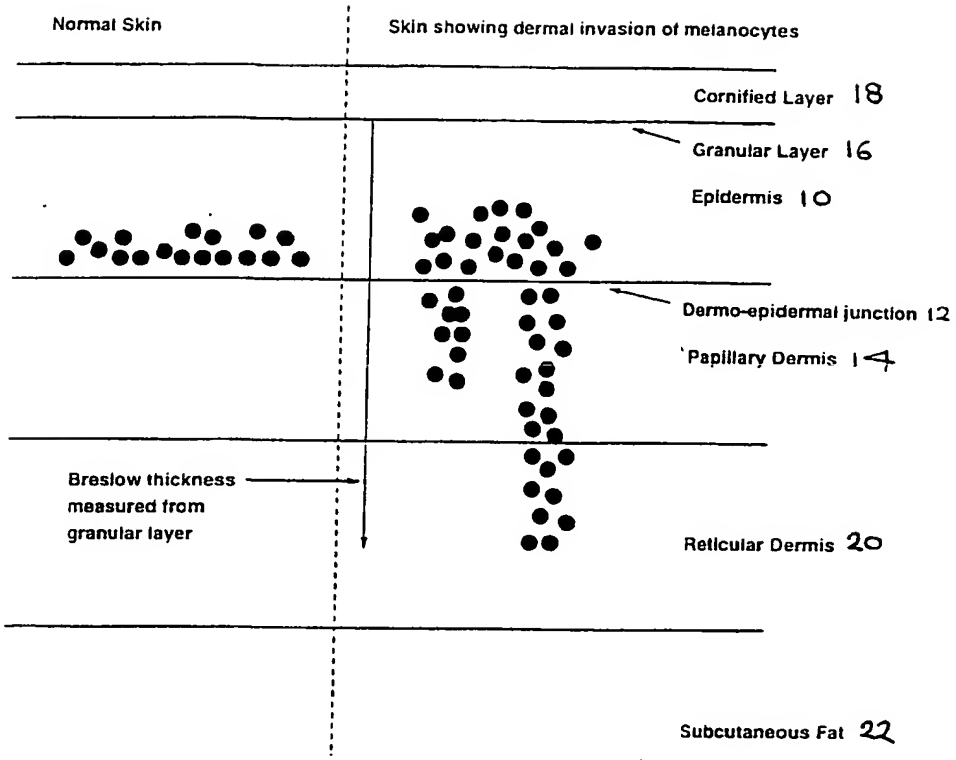
Normal Skin
Skin showing dermal invasion of melanocytes
Cornified Layer 18
Granular Layer 16
Epidermis 10
Dermo-epidermal junction 12
Papillary Dermis 14
Breslow thickness measured from granular layer
Reticular Dermis 20
Subcutaneous Fat 22

FIG. 4

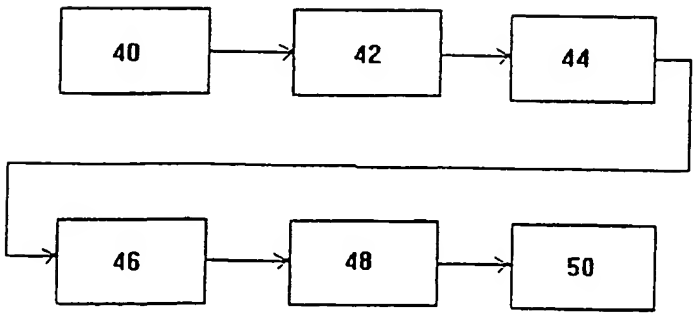
40
42
44
46
48
50

FIG. 5

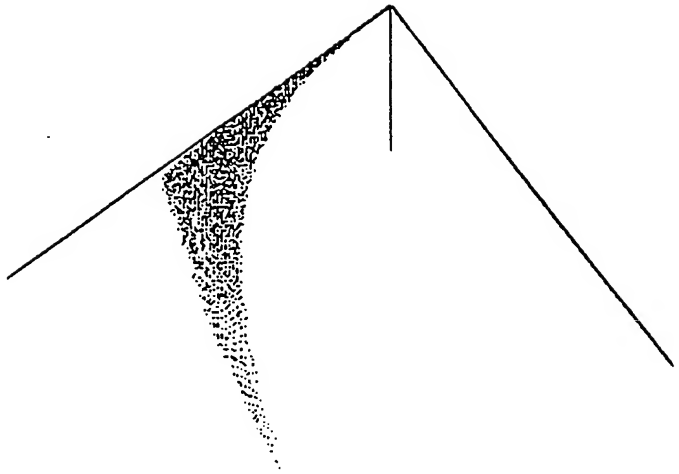

FIG. 6

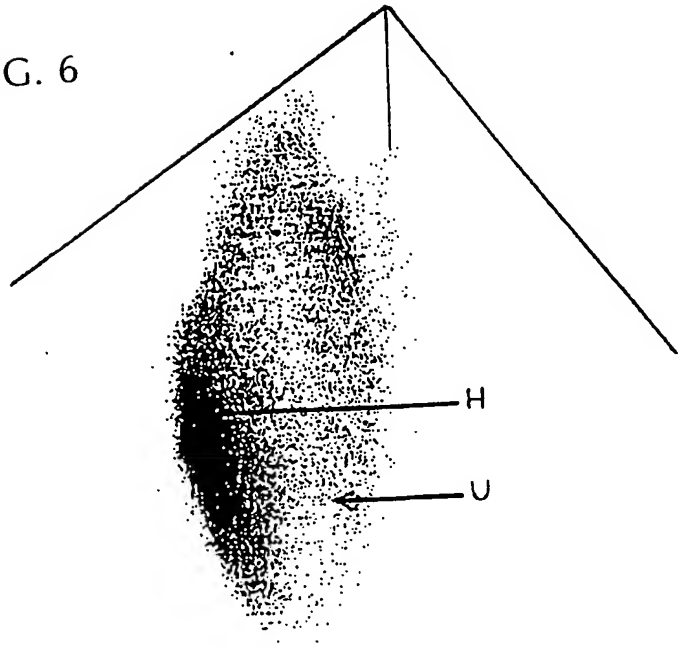

FIG. 7

INTERNATIONAL SEARCH REPORT

Internatic. . Application No
PCT/GB 97/03177

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 A61B5/103

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 A61B G06T

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	JACQUES S L: "THE ROLE OF SKIN OPTICS IN DIAGNOSTIC AND THERAPEUTIC USES OF LASERS" LASERS IN DERMATOLOGY, 1991, pages 1-21, XP000607863 see the whole document ---	
A	WO 90 13091 A (KENET ROBERT O ;KENET BARNEY J (US); TEARNEY GUILLERMO J (US)) 1 November 1990 see abstract; figures ---	
A	US 5 440 388 A (ERICKSON JON W) 8 August 1995 see abstract; figures ---	
	-/--	

[X] Further documents are listed in the continuation of box C.

[X] Patent family members are listed in annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

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Date of the actual completion of the international search	Date of mailing of the international search report
2 March 1998	25.03.98

Name and mailing address of the ISA	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Ferrigno, A

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INTERNATIONAL SEARCH REPORT

International Application No
PCT/GB 97/03177

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	DE 44 27 101 A (BOEHRINGER MANNHEIM GMBH) 1 February 1996 see the whole document ---	
A	US 5 079 698 A (GRENIER LEONARD E ET AL) 7 January 1992 see the whole document ---	
A	WO 96 14795 A (UNIV WASHINGTON) 23 May 1996 see abstract; figures ---	
A	WO 94 16622 A (COMPUTER AIDED MEDICAL INC ;TSIVION YORAM (IL); SHACHAR MENASHE (I) 4 August 1994 see the whole document -----	

2

Form PCT/ISA/210 (continuation of second sheet) (July 1992)

INTERNATIONAL SEARCH REPORT

International application No.
PCT/GB 97/03177

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. [X] Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

see FURTHER INFORMATION sheet PCT/ISA/210

2. [ ] Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:

3. [ ] Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. [ ] As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.

2. [ ] As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. [ ] As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:

4. [ ] No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

[ ] The additional search fees were accompanied by the applicant's protest.

[ ] No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International Application No. PCT/GB 97/03177

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

Claims Nos.: 1 to 14

because they relate to subject matter not required to be searched by this Authority, namely:

Rule 39.1(iv) PCT - Diagnostic method practised on the human or animal body: claims 1-14 relate to a method of detecting abnormal skin tissue as well as to assess the degree of abnormality of said tissue; in particular the claims relate to a method of detecting skin cancer. An incomplete search has been carried out for the invention, to the extent that it is possible in the available documentation.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No
PCT/GB 97/03177

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9013091 A	01-11-90	US 5016173 A	14-05-91
		AU 5524990 A	16-11-90
		US 5241468 A	31-08-93
US 5440388 A	08-08-95	NONE	
DE 4427101 A	01-02-96	AU 3222295 A	04-03-96
		CA 2196187 A	15-02-96
		WO 9604545 A	15-02-96
		EP 0772768 A	14-05-97
US 5079698 A	07-01-92	NONE	
WO 9614795 A	23-05-96	NONE	
WO 9416622 A	04-08-94	AU 6230994 A	15-08-94
		ZA 9400343 A	26-08-94